

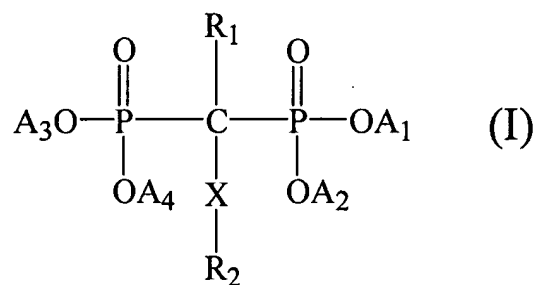
THE PENDING CLAIMS

The following is a complete listing of the pending claims provided as a convenience to the Examiner. No amendments or cancellations are being introduced in this response to the pending election requirement.

LISTING OF CLAIMS

1.-14. (CANCELLED)

15. (PREVIOUSLY PRESENTED) A medicament for treating an autoimmune disease, comprising a treatment enhancing amount of a first active ingredient when in combination with a second active ingredient, wherein the first active ingredient is selected from the group consisting of bisphosphonic acids corresponding to general formula (I)



in which

A₁, A₂, A₃ and A₄ are independently selected from the group consisting of hydrogen, substituted and unsubstituted alkyl, substituted and unsubstituted aryl, substituted and unsubstituted aralkyl, substituted and unsubstituted cycloalkyl, substituted

and unsubstituted heterocyclic residues, metals of Groups I, II and III of the Periodic Table of the elements, and substituted and unsubstituted ammonium or ammonium compounds derived from ethylenediamine or amino acids,

X is absent or is selected from the group consisting of alkylene, alkenylene and hydroxyalkylene,

R₁ and R₂ are independently selected from the group consisting of H, OH, -NH₂, substituted and unsubstituted acyl, substituted and unsubstituted alkyl, substituted and unsubstituted aryl, substituted and unsubstituted cycloalkyl, substituted and unsubstituted aralkyl, substituted and unsubstituted heterocyclic residues, -SR₃, Cl and -NR₃R₄, in which

R₃ and R₄ are independently selected from the group consisting of H, OH, substituted and unsubstituted acyl, substituted and unsubstituted alkyl, substituted and unsubstituted aryl, substituted and unsubstituted aralkyl, substituted and unsubstituted cycloalkyl and substituted and unsubstituted heterocyclic residues, their pharmaceutically compatible salts, esters thereof, salts of the esters and compounds, which upon administration from the compounds according to formula (I) or their salts or esters as metabolites or catabolites,

and a treatment enhancing amount of a second active ingredient when in combination with the first active ingredient, wherein

said second active ingredient is at least one autoantigen specific for the autoimmune disease to be treated and selected from the group consisting of nervous system tissue extracts, collagen, thyroglobulin, acetylcholine receptor protein, DNA, islet

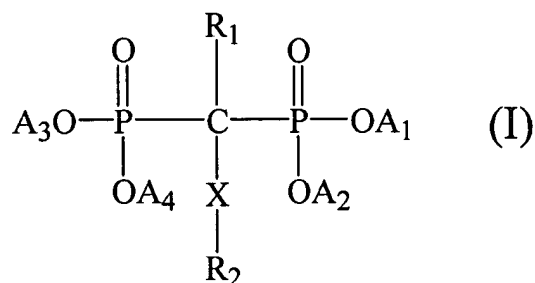
cell extracts, human insulin, liver extracts, adrenal cortex extracts, skin extracts, muscle extracts, haemopoietic cell line extracts, heart extracts, eye lens proteins, S-antigens, gastric cell extracts, parietal cell extracts, intrinsic factor, and intestinal extracts; and an excipient.

16. (PREVIOUSLY PRESENTED) The medicament of claim 15, wherein the bisphosphonic acid is selected from the group consisting of

R_1 is selected from the group consisting of H, OH, $-NH_2$

R_2 is selected from the group consisting of H, OH, $-NH_2$, substituted and unsubstituted acyl, substituted and unsubstituted alkyl having 1 to 12 carbon atoms, substituted and unsubstituted aryl, substituted and unsubstituted cycloalkyl, substituted and unsubstituted aralkyl, substituted and unsubstituted heterocyclic residues, $-SR_3$, C1 and $-NR_3R_4$.

17. (PREVIOUSLY PRESENTED) A medicament for treating an autoimmune disease, comprising
a treatment enhancing amount of a first active ingredient when in combination with a second active ingredient, wherein the first active ingredient is selected from the group consisting of bisphosphonic acids corresponding to general formula (I)



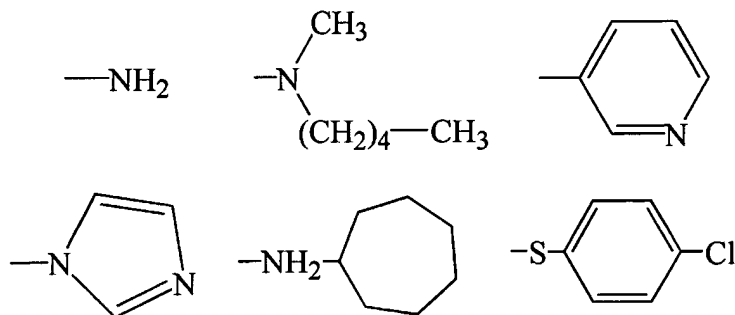
in which

A₁, A₂, A₃ and A₄ are independently selected from the group consisting of hydrogen, substituted and unsubstituted alkyl, substituted and unsubstituted aryl, substituted and unsubstituted aralkyl, substituted and unsubstituted cycloalkyl, substituted and unsubstituted heterocyclic residues, metals of Groups I, II and III of the Periodic Table of the elements, and substituted and unsubstituted ammonium or ammonium compounds derived from ethylenediamine or amino acids,

X is absent or is selected from the group consisting of (CH₂)₁₋₅ and amidino,

R₁ is selected from the group consisting of H and OH, and

R₂ is selected from the group consisting of



their pharmaceutically compatible salts, esters thereof, salts of the esters and compounds, which upon administration form the compounds according to formula (I) or their salts or esters as metabolites or catabolites, and

a treatment enhancing amount of a second active ingredient when in combination with the first active ingredient,

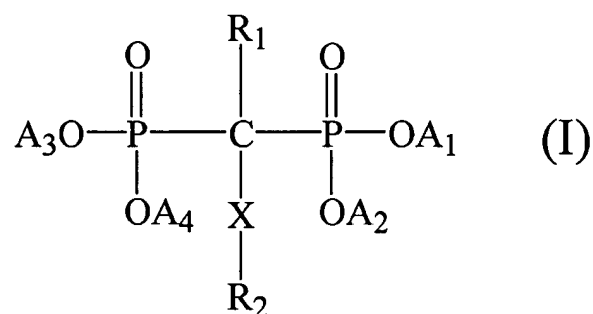
wherein said second active ingredient is at least one autoantigen specific for the autoimmune disease to be treated and selected from the group consisting of nervous system tissue extracts, collagen, thyroglobulin, acetylcholine receptor protein, DNA, islet cell extracts, human insulin, liver extracts, adrenal cortex extracts, skin extracts, muscle extracts, haemopoetic cell line extracts, heart extracts, eye lens proteins, S-antigens, gastric cell extracts, parietal cell extracts, intrinsic factor, and intestinal extracts; and an excipient.

18. (PREVIOUSLY PRESENTED) The medicament of claim 15, wherein the autoantigen is selected from the group consisting of nervous system tissue extracts, islet cell extracts, liver extracts, adrenal cortex extracts, skin extracts, muscle extracts, haemopoetic cell line extracts, heart extracts, gastric cell extracts, parietal cell extracts, and intestinal extracts.
19. (PREVIOUSLY PRESENTED) The medicament of claim 52, wherein the allergen is pollen.
20. (PREVIOUSLY PRESENTED) A medicament according to claim 15 where in the medicament is present in a form selected from the group consisting of solid form, ointment, solution, and spray.

21.-51. (CANCELLED)

52. (PREVIOUSLY PRESENTED) A medicament for treating an allergy, comprising:
a treatment enhancing amount of a first active ingredient when in combination with a
second active ingredient, wherein the first active ingredient is selected from the group
consisting of

bisphosphonic acids corresponding to general formula (I)



in which

A₁, A₂, A₃ and A₄ are independently selected from the group consisting of
hydrogen, substituted and unsubstituted alkyl, substituted and unsubstituted aryl,
substituted and unsubstituted aralkyl, substituted and unsubstituted cycloalkyl, substituted
and unsubstituted heterocyclic residues, metals of Groups I, II and III of the Periodic
Table of the elements, and substituted and unsubstituted ammonium or ammonium
compounds derived from ethylenediamine or amino acids,

X is absent or is selected from the group consisting of alkylene, alkenylene and
hydroxyalkylene,

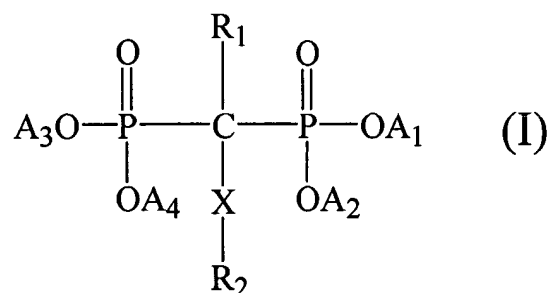
R_1 and R_2 are independently selected from the group consisting of H, OH, $-NH_2$, substituted and unsubstituted acyl, substituted and unsubstituted alkyl, substituted and unsubstituted aryl, substituted and unsubstituted cycloalkyl, substituted and unsubstituted aralkyl, substituted and unsubstituted heterocyclic residues, $-SR_3$, Cl and $-NR_3R_4$, in which

R_3 and R_4 are independently selected from the group consisting of H, OH, substituted and unsubstituted acyl, substituted and unsubstituted alkyl, substituted and unsubstituted aryl, substituted and unsubstituted aralkyl, substituted and unsubstituted cycloalkyl and substituted and unsubstituted heterocyclic residues, their pharmaceutically compatible salts, esters thereof, salts of the esters and compounds, which upon administration from the compounds according to formula (I) or their salts or esters as metabolites or catabolites,

and a treatment enhancing amount of a second active ingredient when in combination with the first active ingredient, wherein said second active ingredient is an allergen ingredient specific for the allergy to be treated and is selected from the group consisting of pollen, nickel, and food; and

an excipient.

53. (PREVIOUSLY PRESENTED) A medicament for treating an allergy, comprising:
a treatment enhancing amount of a first active ingredient when in combination with a second active ingredient, wherein the first active ingredient is selected from the group consisting of bisphosphonic acids corresponding to general formula (I)



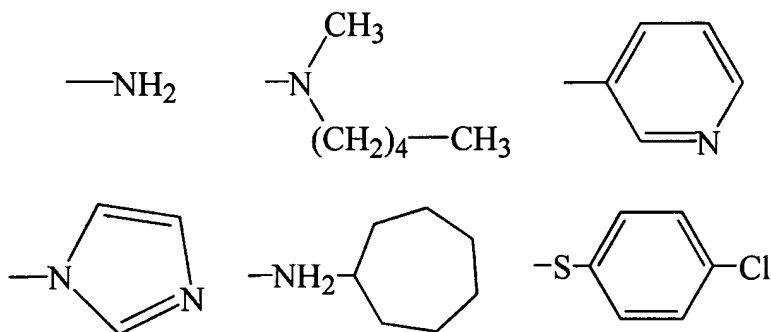
in which

A₁, A₂, A₃ and A₄ are independently selected from the group consisting of hydrogen, substituted and unsubstituted alkyl, substituted and unsubstituted aryl, substituted and unsubstituted aralkyl, substituted and unsubstituted cycloalkyl, substituted and unsubstituted heterocyclic residues, metals of Groups I, II and III of the Periodic Table of the elements, and substituted and unsubstituted ammonium or ammonium compounds derived from ethylenediamine or amino acids,

X is absent or is selected from the group consisting of (CH₂)₁₋₅ and amidino,

R₁ is selected from the group consisting of H and OH, and

R₂ is selected from the group consisting of



their pharmaceutically compatible salts, esters thereof, salts of the esters and compounds, which upon administration form the compounds according to formula (I) or their salts or esters as metabolites or catabolites, and

a treatment enhancing amount of a second active ingredient when in combination with the first active ingredient, wherein said second active ingredient is an allergen ingredient specific for the allergy to be treated and selected from the group consisting of pollen, nickel, and food; and
an excipient.

54. (PREVIOUSLY PRESENTED) The medicament of claim 15, wherein the autoantigen is from a nervous system tissue extract and the autoantigen is myelin basic protein.
55. (PREVIOUSLY PRESENTED) The medicament of claim 15, wherein the autoantigen is selected from the group consisting of collagen, thyroglobulin, acetylcholine receptor protein, human insulin, eye lens proteins, S-antigens, and intrinsic factor.
56. (PREVIOUSLY PRESENTED) The medicament of claim 15, wherein the autoantigen is DNA.
57. (PREVIOUSLY PRESENTED) The medicament of Claim 52, wherein the allergen ingredient is food.
58. (PREVIOUSLY PRESENTED) The medicament of Claim 15 wherein the bisphosphonic acid is an amino-1-hydroxyalkylidene-1,1-bisphosphonic acid wherein the alkyl is methyl, ethyl, propyl, butyl, or hexyl.

59. (PREVIOUSLY PRESENTED) The medicament of Claim 15 wherein the bisphosphonic acid is amidinomethylenebisphosphonic acid, risedronic acid, zoledronic acid, cimidronic acid, or tiludronic acid.
60. (PREVIOUSLY PRESENTED) The medicament of Claim 52 wherein the bisphosphonic acid is an amino-l-hydroxyalkylidene-1,1-bisphosphonic acid wherein the alkyl is methyl, ethyl, propyl, butyl, or hexyl.
61. (PREVIOUSLY PRESENTED) The medicament of Claim 52 wherein the bisphosphonic acid is amidinomethylenebisphosphonic acid, risedronic acid, zoledronic acid, cimidronic acid, or tiludronic acid.
62. (PREVIOUSLY PRESENTED) The medicament of Claim 15 wherein the bisphosphonic acid is an amino-l-hydroxyalkylidene-1,1-bisphosphonic acid wherein the alkyl is methyl, ethyl, propyl, butyl, or hexyl; and the autoantigen is from a nervous system tissue extract and is myelin basic protein.
63. (PREVIOUSLY PRESENTED) The medicament of Claim 15 wherein the bisphosphonic acid is an amino-l-hydroxyalkylidene-1,1-bisphosphonic acid wherein the alkyl is methyl, ethyl, propyl, butyl, or hexyl; and the autoantigen is collagen.

64. (PREVIOUSLY PRESENTED) The medicament of Claim 15 wherein the bisphosphonic acid is an amino-1-hydroxyalkylidene-1,1-bisphosphonic acid wherein the alkyl is methyl, ethyl, propyl, butyl, or hexyl; and the autoantigen is insulin,
65. (PREVIOUSLY PRESENTED) The medicament of Claim 15 wherein the bisphosphonic acid and the autoantigen are present in a form for separate administration.

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